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A Commemorative Issue in Honor of Professor Valder R. Arruda: Hemophilia and Rare Bleeding Disorders

Guest Editor:

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Deadline for manuscript submissions:
closed (31 August 2022)

Message from the Guest Editor

Dear Colleagues,

Rare bleeding disorders (RBD) and hemophilia are inherited coagulopathies with variable bleeding phenotypes, whose hemostatic control was, until recently, based upon replacement therapy. In patients with bleeding symptoms, laboratory assessment and, especially, molecular workup enable an accurate diagnosis, as well as prenatal and family counseling. In advance phase studies on gene therapy for hemophilia, the control of critical safety and efficacy challenges will pave the way for approval as well as to set up the basis for future gene-based strategies for RBD. In this Special Issue, Molecular and Functional Research in Hemophilia and Rare Bleeding Disorders, we aim to discuss some of these important aspects associated with diagnosis and potential care.

Prof. Dr. Gili Kenet
Dr. Valder R. Arruda
Guest Editors

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Message from the Editor-in-Chief

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